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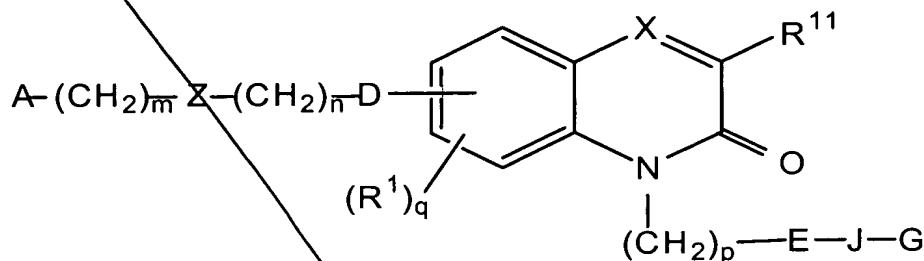
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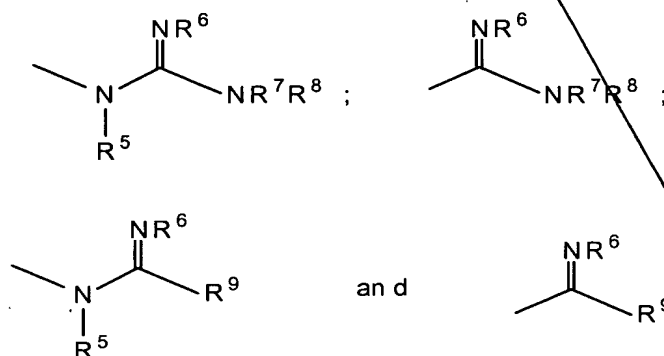
WHAT IS CLAIMED IS:

1. A compound of having the following formula:



wherein:

A is a member selected from the group consisting of: R^2 , $-NR^3R^4$, $-C(=O)NR^3R^4$,



where R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , and R^9 are independently selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S; where R^6 taken with either of R^7 and R^8 , and/or R^7 taken with R^8 , can each form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of

N, O and S;

m is an integer from 0-3;

Q!
Cont 5 Z is a member selected from the group consisting of a direct link, C₁₋₈alkyl, C₃₋₈cycloalkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₁₋₈carbocyclic aryl, or a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S;

n is an integer from 0-3;

10 D is a member selected from the group consisting of a direct link, -CH₂-, -O-, -N(R²)-, -C(=O)-, -S-, -SO₂-, -SO₂-N(R²)-, -N(R²)-SO₂-, -OC(=O)-, -C(=O)O-, -C(=O)-N(R²)- and -N(R²)-C(=O)-;

15 R¹ is a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, halogen, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH, C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, -CN, -NO₂, C₀₋₈alkyl-OH, C₀₋₈alkyl-SH, -C(=O)NR²R³, -O-R² and -O-C(=O)R², an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, polyhaloalkyl, -SO₂R², C₀₋₈alkyl-C(=O)OH and C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, where R² and R³ is as described above;

20

q is an integer from 0-3;

X is N or -CR¹²;

25 R¹¹ and R¹² are independently a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, C₆₋₁₂carbocyclic aryl, C₁₋₆alkylaryl, C₁₋₆alkyl-C₃₋₈cycloalkyl, -O-R², -O-C(=O)R², -C₁₋₈alkyl-O-R¹⁰, -C₁₋₈alkyl-O-C(=O)R¹⁰, -C₁₋₈alkyl-C(=O)OR¹⁰, -C₁₋₈alkyl-O-C(=O)OR¹⁰, -C₁₋₈alkyl-C(=O)NR¹⁰R¹⁰, -C₁₋₈alkyl-NR¹⁰R¹⁰, -C₁₋₈alkyl-NR¹⁰C(=O)R¹⁰, -SR¹⁰, where R² is as described above and R¹⁰ is a member selected from the group consisting of H,

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cont 5 C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, and wherein when two R¹⁰ groups are present they may be taken together to form a saturated or unsaturated ring with the atom to which they are both attached;

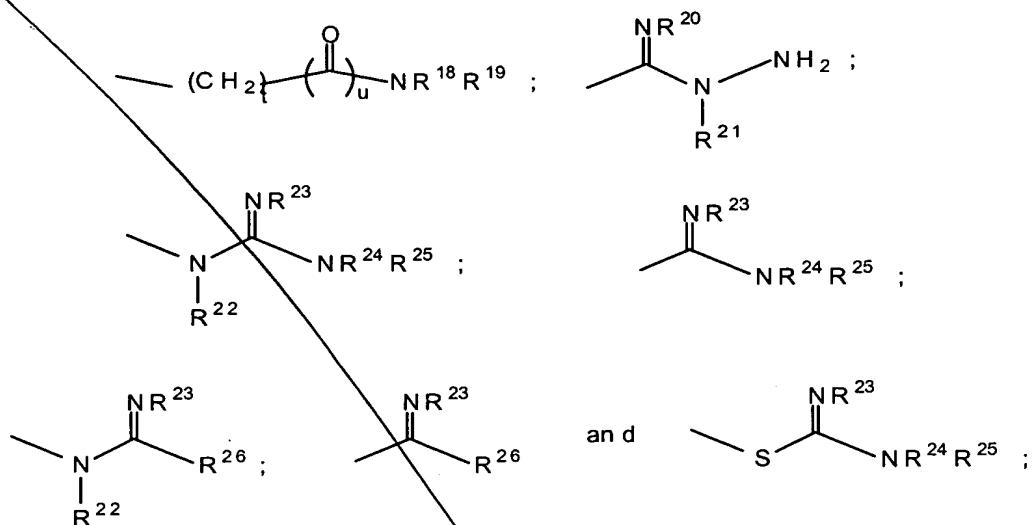
p is an integer from 0-3;

E is a member selected from the group consisting of a direct link, -O-, -N(-R¹¹)-, where R¹¹ is as set forth above, phenylene, a bivalent 5 to 12 member heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S, wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by from 0 to 5 R¹⁴ groups;

J is a member selected from the group consisting of a direct link, a bivalent C₃₋₈cycloalkyl group, phenylene, a 5 to 12 member bivalent heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by from 0 to 5 R¹⁴ groups;

each R¹⁴ group is a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, halogen, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH, C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, -CN, -NO₂, C₀₋₈alkyl-OH, C₀₋₈alkyl-SH, -O-R² and -O-C(=O)R², an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH and C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl;

G is a member selected from the group consisting of: H; -CN; -OR¹⁷;



wherein

t is an integer from 0 to 6,

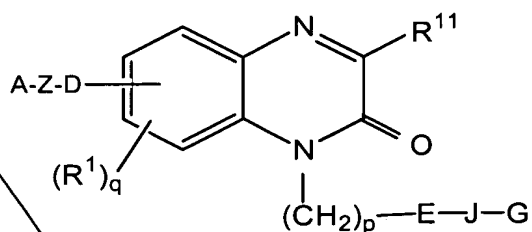
- u is the integer 0 or 1, and R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵ and R²⁶ are
- 5 independently selected from the group consisting of H, -OH, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, C₆₋₁₂carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C₁₋₆alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S;
- 10 where R¹⁸ taken with R¹⁹, R²² taken with either of R²⁴ and R²⁵, and R²⁴ taken with R²⁵, can each independently form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

with the proviso that when G is H, -CN, -OR¹⁷, either E or J must contain at least one N atom;

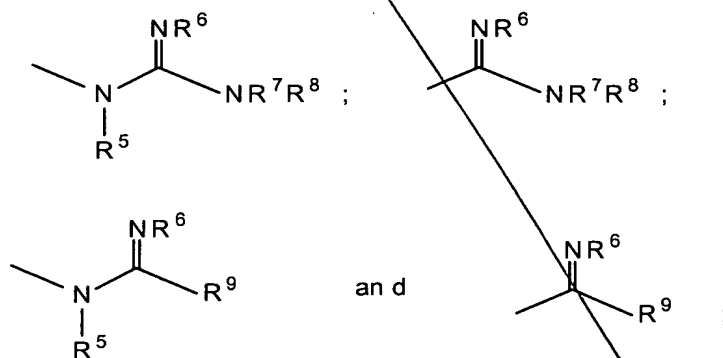
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and all pharmaceutically acceptable isomers, salts, hydrates, solvates and prodrug derivatives thereof.

2. A compound of formula II:



5 A is a member selected from the group consisting of: R^2 , $\text{-NR}^3\text{R}^4$, $\text{-C(=O)NR}^3\text{R}^4$,



10 where R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , and R^9 are independently selected from the group consisting of H, -OH , $\text{C}_{1-8}\text{alkyl}$, $\text{C}_{2-8}\text{alkenyl}$, $\text{C}_{2-8}\text{alkynyl}$, $\text{C}_{3-8}\text{cycloalkyl}$, $\text{C}_{6-12}\text{carbocyclic aryl}$, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and $\text{C}_{1-6}\text{alkylheterocyclic ring system}$ having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S; where R^6 taken with either of R^7 and R^8 , and/or R^7 taken with R^8 , can each form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

Z is a member selected from the group consisting of a direct link, $\text{C}_{1-8}\text{alkyl}$,

Q1 cont
C₃₋₈cycloalkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₁₋₈carbocyclic aryl, or a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S;

D is a member selected from the group consisting of a direct link, -CH₂-, -O-, -N(R²)-, -C(=O)-, -S-, -SO₂-, -SO₂-N(R²)-, -N(R²)-SO₂-, -OC(=O)-, -C(=O)O-, -C(=O)-N(R²)-, and -N(R²)-C(=O)-;

R¹ is a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, halogen, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH, C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, -CN, -NO₂, C₀₋₈alkyl-OH, C₀₋₈alkyl-SH, -C(=O)NR²R³, -O-R² and -O-C(=O)R², an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, polyhaloalkyl, -SO₂R², C₀₋₈alkyl-C(=O)OH and C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, where R² and R³ is as described above;

q is an integer from 0-3;

R¹¹ is independently a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, C₆₋₁₂carbocyclic aryl, C₁₋₆alkylaryl, C₁₋₆alkyl-C₃₋₈cycloalkyl, -O-R², -O-C(=O)R², -C₁₋₈alkyl-O-R¹⁰, -C₁₋₈alkyl-O-C(=O)R¹⁰, -C₁₋₈alkyl-C(=O)OR¹⁰, -C₁₋₈alkyl-O-C(=O)OR¹⁰, -C₁₋₈alkyl-C(=O)NR¹⁰R¹⁰, -C₁₋₈alkyl-NR¹⁰R¹⁰, -C₁₋₈alkyl-NR¹⁰C(=O)R¹⁰, -SR¹⁰, where R² is as described above and R¹⁰ is a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, and wherein when two R¹⁰ groups are present they may be taken together to form a saturated or unsaturated ring with the atom to which they are both attached;

p is an integer from 0-2;

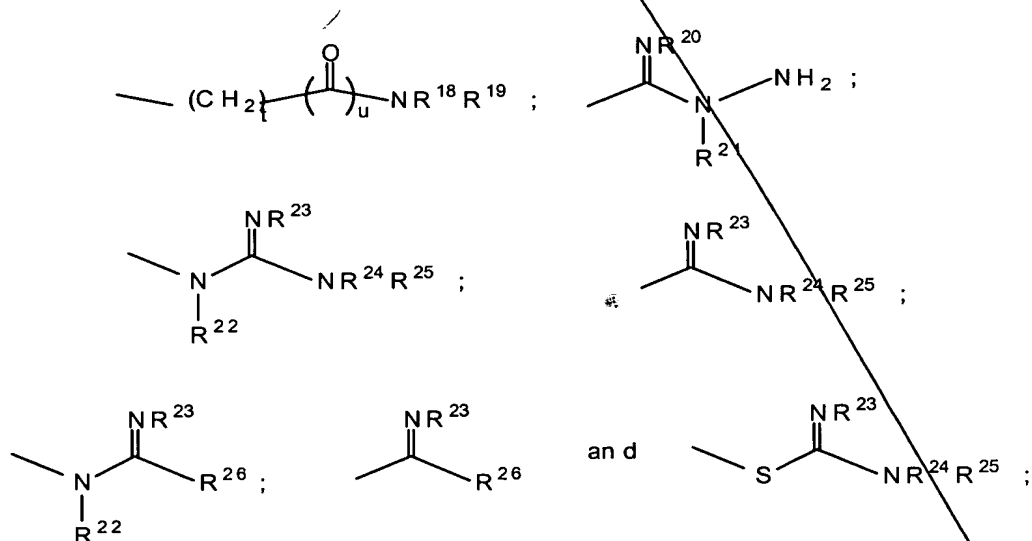
E is a member selected from the group consisting of a direct link, -O-, -N(-R¹¹)-, where R¹¹ is as set forth above, phenylene, a bivalent 5 to 12 member heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of

N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S, wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by from 0 to 5 R¹⁴ groups;

5 J is a member selected from the group consisting of a direct link, a bivalent C₃₋₈cycloalkyl group, phenylene, a 5 to 12 member bivalent heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by
10 from 0 to 5 R¹⁴ groups;

each R¹⁴ group is a member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, halogen, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH, C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl, -CN, -NO₂, C₀₋₈alkyl-OH, C₀₋₈alkyl-SH, -O-R² and
15 -O-C(=O)R², an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH and C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl;

G is a member selected from the group consisting of: H; -CN; -OR¹⁷;



wherein

t is an integer from 0 to 6,

u is the integer 0 or 1, and R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{24} , R^{25} and R^{26} are independently selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S; where R^{18} taken with R^{19} , R^{22} taken with either of R^{24} and R^{25} , and R^{24} taken with R^{25} , can each independently form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

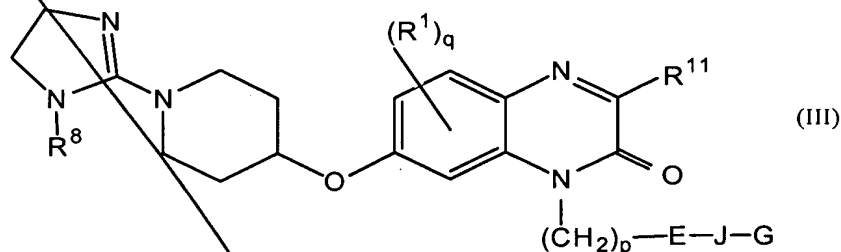
with the proviso that when G is H, -CN, -OR¹⁷, either E or J must contain at least one N atom;

and all pharmaceutically acceptable isomers, salts, hydrates, solvates and prodrug derivatives thereof.

3. A compound of claim 2, wherein D is a member selected from the group consisting of: -O-, -NR²-, -C(=O)-, -S-, -SO₂-, -SO₂-NR²-, -NR²-SO₂-, -OC(=O)-, -C(=O)NR², and -NR²-C(=O)-.

4. A compound of claim 3, wherein D is a member selected from the group consisting of: -O-, -NR²-, -C(=O)-, -S-, and -SO₂-.

5. A compound of formula III:



5 wherein:

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R^8 is selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S;

R^1 is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, halogen, polyhaloalkyl, C_{0-8} alkyl-C(=O)OH, C_{0-8} alkyl-C(=O)O- C_{1-8} alkyl, -CN, -NO₂, C_{0-8} alkyl-OH, C_{0-8} alkyl-SH, -C(=O)NR²R³, -O-R² and -O-C(=O)R², an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, polyhaloalkyl, -SO₂R², C_{0-8} alkyl-C(=O)OH and C_{0-8} alkyl-C(=O)O- C_{1-8} alkyl, where R² and R³ is as described above;

R^2 is selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S;

25 q is 0-3;

Q² cont 5 R^{11} is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, C_{1-6} alkylaryl, C_{1-6} alkyl- C_{3-8} cycloalkyl, $-O-R^2$, $-O-C(=O)R^2$, $-C_{1-8}$ alkyl- $O-R^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)R^{10}$, $-C_{1-8}$ alkyl- $C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $C(=O)NR^{10}R^{10}$, $-C_{1-8}$ alkyl- $NR^{10}R^{10}$, $-C_{1-8}$ alkyl- $NR^{10}C(=O)R^{10}$, $-SR^{10}$, where R^2 is as described above and R^{10} is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, and wherein when two R^{10} groups are present they may be taken together to form a saturated or unsaturated ring with the atom to which they are both attached;

10 p is an integer from 0-2;

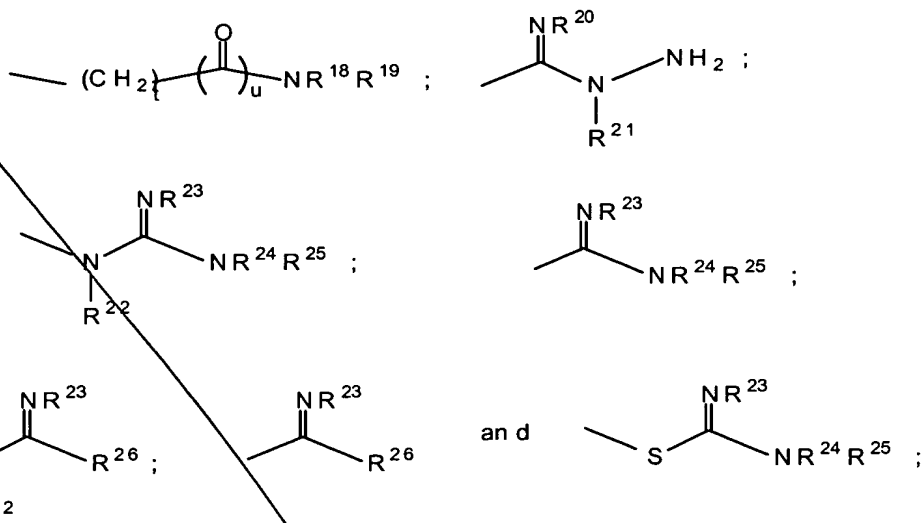
E is a member selected from the group consisting of a direct link, $-O-$, $-N(R^{11})-$, where R^{11} is as set forth above, phenylene, a bivalent 5 to 12 member heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S, wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by from 0 to 5 R^{14} groups;

J is a member selected from the group consisting of a direct link, a bivalent C_{3-8} cycloalkyl group, phenylene, a 5 to 12 member bivalent heteroaryl group containing 1 to 4 heteroatoms selected from the group consisting of N, O and S, and a five to ten membered non-aromatic bivalent heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S wherein said heteroaryl and said non-aromatic heterocyclic ring structure may be independently substituted by from 0 to 5 R^{14} groups;

25 each R^{14} group is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, halogen, polyhaloalkyl, C_{0-8} alkyl- $C(=O)OH$, C_{0-8} alkyl- $C(=O)O-C_{1-8}$ alkyl, $-CN$, $-NO_2$, C_{0-8} alkyl- OH , C_{0-8} alkyl- SH , $-O-R^2$ and $-O-C(=O)R^2$, an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl,

C_{3-8} cycloalkyl, polyhaloalkyl, C_{0-8} alkyl-C(=O)OH and C_{0-8} alkyl-C(=O)O- C_{1-8} alkyl;

G is a member selected from the group consisting of: H; -CN; -OR¹⁷;



wherein

t is an integer from 0 to 6,

- 5 u is the integer 0 or 1, and R¹⁷, R¹⁸, R¹⁹, R²⁰, R²¹, R²², R²³, R²⁴, R²⁵ and R²⁶ are independently selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S;
- 10 where R¹⁸ taken with R¹⁹, R²² taken with either of R²⁴ and R²⁵, and R²⁴ taken with R²⁵, can each independently form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

- with the proviso that when G is H, -CN, -OR¹⁷, either E or J must contain at
- 15 least one N atom;

and all pharmaceutically acceptable isomers, salts, hydrates, solvates and prodrug derivatives thereof.

6. A compound of claim 5, wherein R^1 and R^8 are independently a lower alkyl group and R^{11} is hydrogen or is a C_1 to C_8 alkyl group.

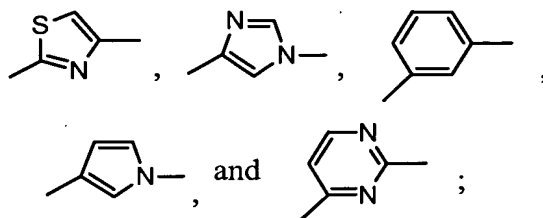
7. A compound of claim 5, wherein q is zero and R^8 is lower alkyl group.

8. A compound of claim 5, wherein:

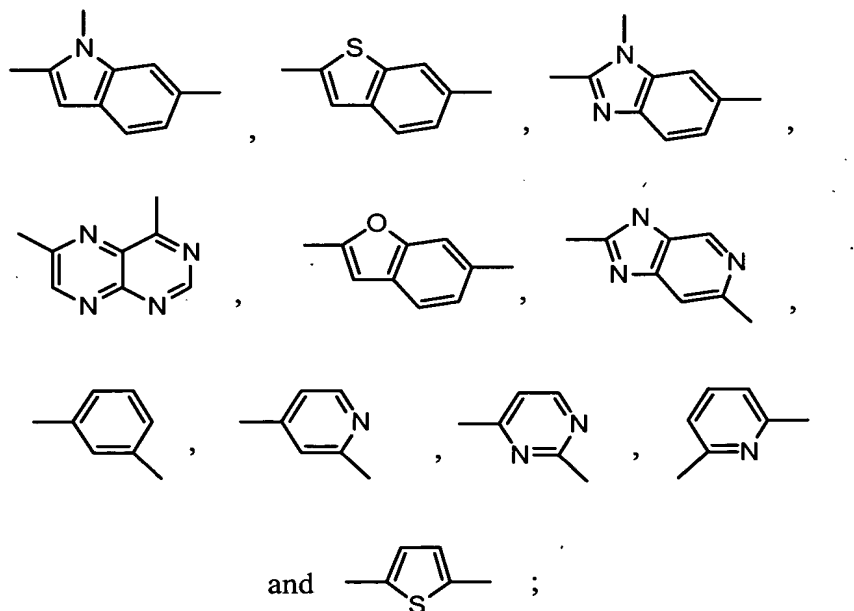
5 R^8 is a methyl group;

p is an integer from 1-2;

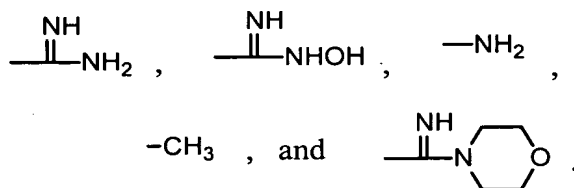
E is selected from the group consisting of: a direct link,



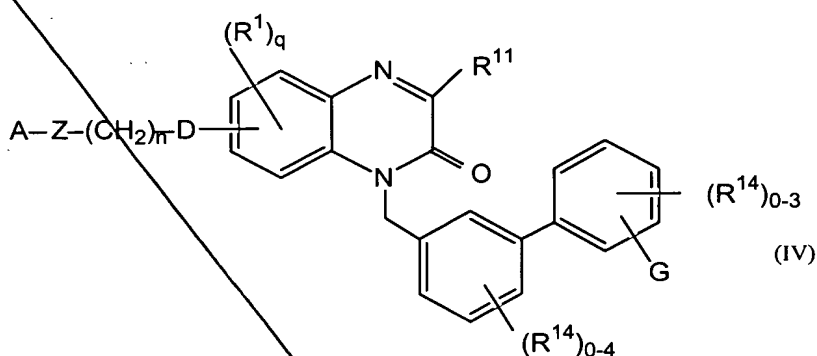
J is selected from the group consisting of:



and G is selected from the group consisting of:

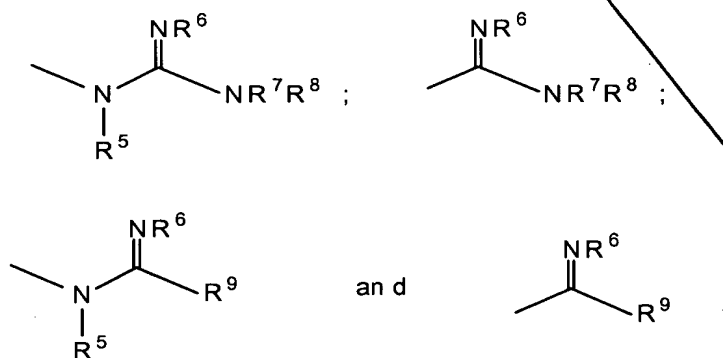


9. A compound of formula IV:



wherein:

5 A is a member selected from the group consisting of: R^2 , ---NR^3R^4 , ---C(=O)NR^3R^4 ,



10 where R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , and R^9 are independently selected from the group consisting of H, ---OH , $\text{C}_{1-8}\text{alkyl}$, $\text{C}_{2-8}\text{alkenyl}$, $\text{C}_{2-8}\text{alkynyl}$, $\text{C}_{3-8}\text{cycloalkyl}$,

Q3
cont
5 C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and

C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S; where r^6 taken with either of R^7 and R^8 , and/or R^7 taken with R^8 , can each form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

10 Z is a member selected from the group consisting of a direct link, C_{1-8} alkyl, C_{3-8} cycloalkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{1-8} carbocyclic aryl, or a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S;

n is 0-3;

15 D is a member selected from the group consisting of: $-CH_2-$, $-O-$, $-N R^2$, $-C(=O)-$, $-S-$, $-SO_2-$, $-SO_2-NR^2$, $-NR^2-SO_2$, $-OC(=O)-$, $-C(=O)NR^2$, and $-NR^2-C(=O)-$;

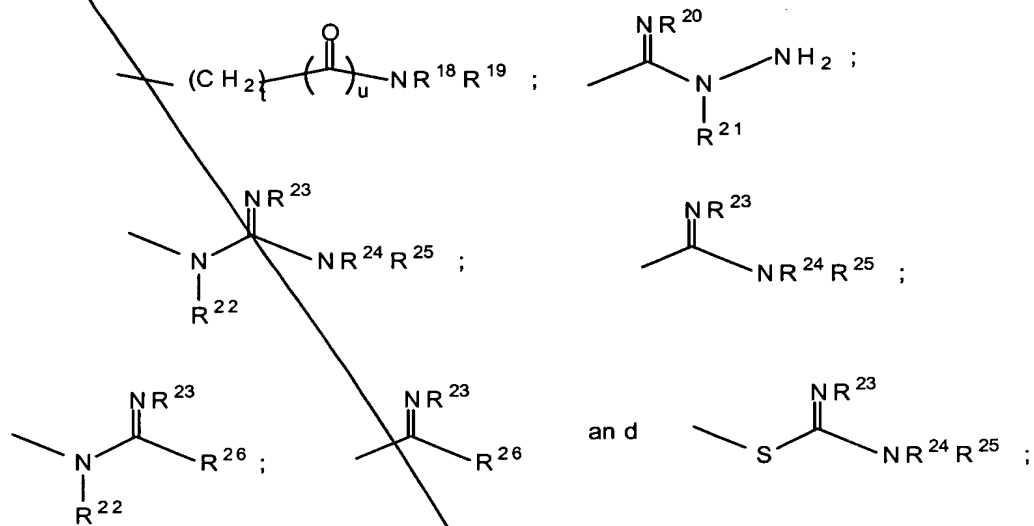
20 R^1 and R^{14} are independently a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, halogen, polyhaloalkyl, C_{0-8} alkyl- $C(=O)OH$, C_{0-8} alkyl- $C(=O)O-C_{1-8}$ alkyl, $-CN$, $-NO_2$, C_{0-8} alkyl-OH, C_{0-8} alkyl-SH, $-O-R^2$ and $-O-C(=O)R^2$, an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, polyhaloalkyl, C_{0-8} alkyl- $C(=O)OH$ and C_{0-8} alkyl- $C(=O)O-C_{1-8}$ alkyl;

25 q is 0-3;

30 R^{11} is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, C_{1-6} alkylaryl, C_{1-6} alkyl- C_{3-8} cycloalkyl, $-O-R^2$, $-O-C(=O)R^2$, $-C_{1-8}$ alkyl- $O-R^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)R^{10}$, $-C_{1-8}$ alkyl- $C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $C(=O)NR^{10}R^{10}$,

$-C_{1-8}\text{alkyl}-NR^{10}R^{10}$, $-C_{1-8}\text{alkyl}-NR^{10}C(=O)R^{10}$, $-SR^{10}$, where R^2 is as described above and R^{10} is a member selected from the group consisting of H, $C_{1-8}\text{alkyl}$, $C_{2-8}\text{alkenyl}$, $C_{2-8}\text{alkynyl}$, and wherein when two R^{10} groups are present they may be taken together to form a saturated or unsaturated ring with the atom to which they are both attached;

5 G is a member selected from the group consisting of: H; $-CN$; $-OR^{17}$;



wherein

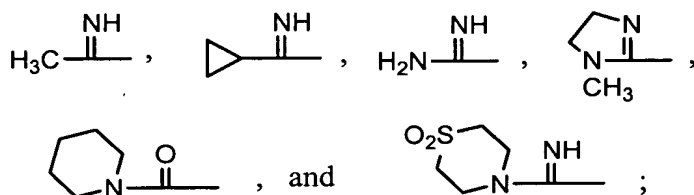
t is an integer from 0 to 6,

u is the integer 0 or 1, and R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{24} , R^{25} and R^{26} are independently selected from the group consisting of H, $-OH$, $C_{1-8}\text{alkyl}$, $C_{2-8}\text{alkenyl}$, $C_{2-8}\text{alkynyl}$, $C_{3-8}\text{cycloalkyl}$, $C_{6-12}\text{carbocyclic aryl}$, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and $C_{1-6}\text{alkylheterocyclic ring system}$ having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S; where R^{18} taken with R^{19} , R^{22} taken with either of R^{24} and R^{25} , and R^{24} taken with R^{25} ,
10
15 can each independently form a 5 to 6 membered heterocyclic ring containing from 1 to 4 atoms selected from the group consisting of N, O and S;

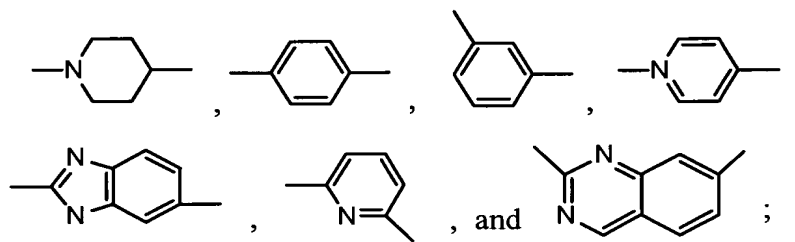
with the proviso that when G is H, $-CN$, $-OR^{17}$, either E or J must contain at least one N atom;

and all pharmaceutically acceptable isomers, salts, hydrates, solvates and prodrug derivatives thereof.

10. A compound of claim 9, wherein R^1 , R^8 , R^{11} and R^{14} are independently selected from the group consisting of hydrogen, methyl and ethyl;
A is selected from the group consisting of: -H, -CH₃, -NH₂, -C(O)N(CH₃)₂,



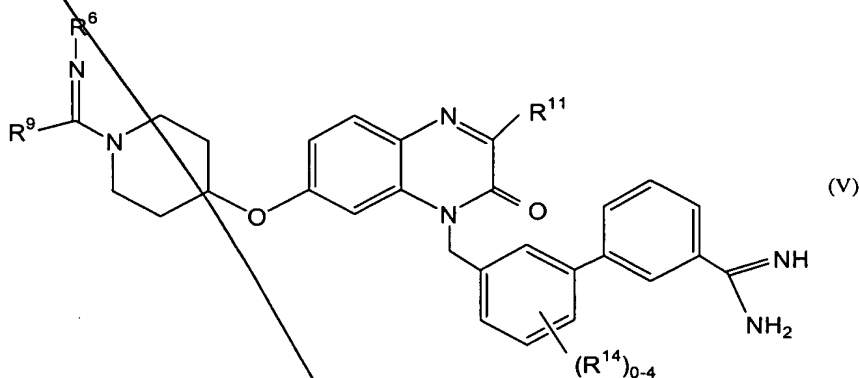
Z is selected from the group consisting of:



n is an integer from 0-2; and

D is selected from the group consisting of: -O-, -N(CH₃)-, and -CH₂-.

11. A compound of formula V:



wherein:

R^2 , R^6 , and R^9 are independently selected from the group consisting of H, -OH, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, a five to ten membered heterocyclic ring system containing 1-4 heteroatoms selected from the group consisting of N, O and S; and C_{1-6} alkylheterocyclic ring system having in the ring system 5 to 10 atoms with 1 to 4 of such atoms being selected from the group consisting of N, O and S;

R^{11} is independently a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, C_{6-12} carbocyclic aryl, C_{1-6} alkylaryl, C_{1-6} alkyl- C_{3-8} cycloalkyl, $-O-R^2$, $-O-C(=O)R^2$, $-C_{1-8}$ alkyl- $O-R^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)R^{10}$, $-C_{1-8}$ alkyl- $C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $O-C(=O)OR^{10}$, $-C_{1-8}$ alkyl- $C(=O)NR^{10}R^{10}$, $-C_{1-8}$ alkyl- $NR^{10}R^{10}$, $-C_{1-8}$ alkyl- $NR^{10}C(=O)R^{10}$, $-SR^{10}$, where R^2 is as described above and R^{10} is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, and wherein when two R^{10} groups are present they may be taken together to form a saturated or unsaturated ring with the atom to which they are both attached;

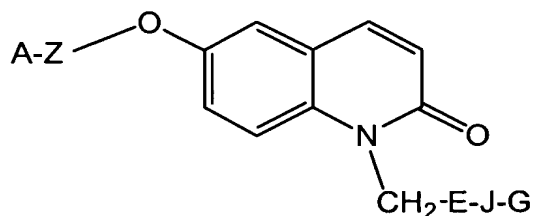
each R^{14} group is a member selected from the group consisting of H, C_{1-8} alkyl, C_{2-8} alkenyl, C_{2-8} alkynyl, C_{3-8} cycloalkyl, halogen, polyhaloalkyl, C_{0-8} alkyl- $C(=O)OH$, C_{0-8} alkyl- $C(=O)O-C_{1-8}$ alkyl, -CN, $-NO_2$, C_{0-8} alkyl-OH, C_{0-8} alkyl-SH, $-O-R^2$ and $-O-C(=O)R^2$, an unsubstituted amino group, a mono- or di-substituted amino group, wherein the substituted amino groups are independently substituted by at least one

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cont

member selected from the group consisting of H, C₁₋₈alkyl, C₂₋₈alkenyl, C₂₋₈alkynyl, C₃₋₈cycloalkyl, polyhaloalkyl, C₀₋₈alkyl-C(=O)OH and C₀₋₈alkyl-C(=O)O-C₁₋₈alkyl;

and all pharmaceutically acceptable isomers, salts, hydrates, solvates and
5 prodrug derivatives thereof.

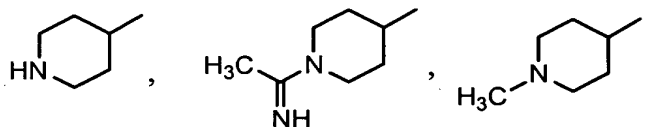
12. A compound having the following structure:



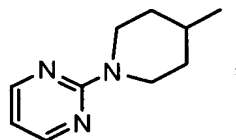
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wherein:

A-Z is a member selected from the group consisting of:

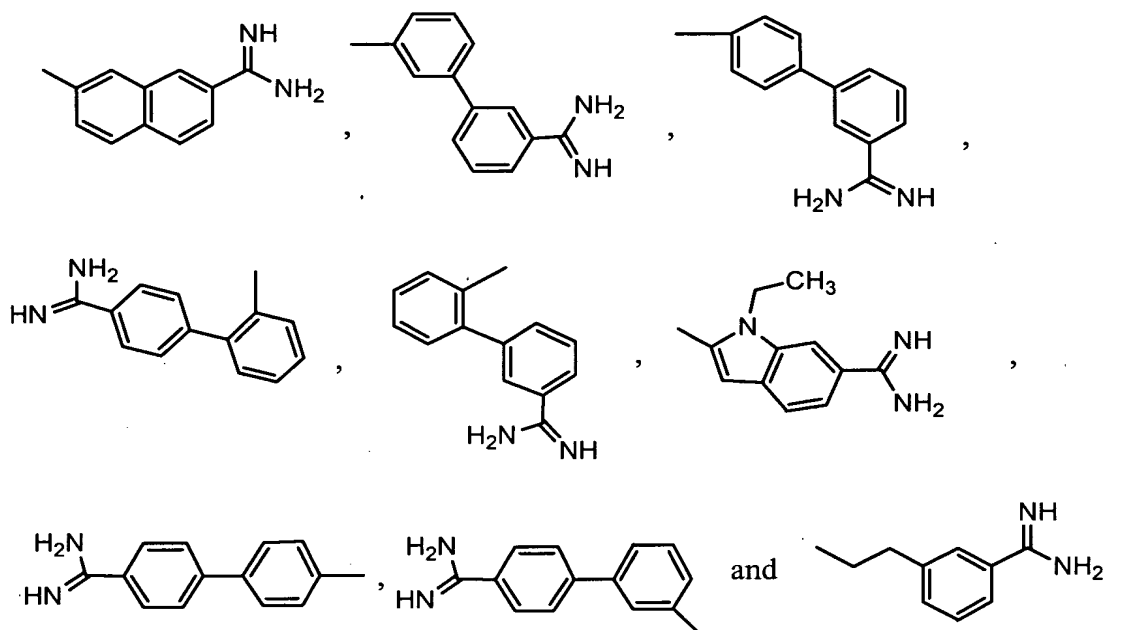


and



15

E-J-G is a member selected from the group consisting of:



and all pharmaceutically acceptable isomers, salts, hydrates, solvates and prodrug derivatives thereof.

5

13. A pharmaceutical composition for preventing or treating a condition in a mammal characterized by undesired thrombosis comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound as in one of claims 1-12.

10

14. A method for preventing or treating a condition in a mammal characterized by undesired thrombosis comprising administering to said mammal a therapeutically effective amount of a compound as in one of claims 1-12.

15

15. The method of claim 14, wherein the condition is selected from the group consisting of:

acute coronary syndrome, myocardial infarction, unstable angina, refractory angina, occlusive coronary thrombus occurring post-thrombolytic therapy or post-

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16. A method for inhibiting the coagulation of biological samples comprising the administration of a compound as in one of claims 1-12.